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INTERNATIONAL PRELIMINARY EXAMINATION REPORT

(PCT Article 36 and Rule 70)

	Applicant's or agent's file reference								
5-70137A			FOR FURTHER ACTION	See Notification of Transmittal of International Preliminary Examination Report (Form PCT/IPEA/416)					
International application No.			International filing date (day/mon 09.10.2003	10.10.2002					
International Patent Classification (IPC) or both national classification and IPC C07C259/06 Applicant									
	Applicant -YNGENTA PARTICIPATIONS AG et al								
1,	This international preliminary examination report has been prepared by this international Preliminary Examining Authority and is transmitted to the applicant according to Article 36.								
2.	This REPORT consists of a total of 5 sheets, including this cover sheet.								
	This report is also accompanied by ANNEXES, i.e. sheets of the description, claims and/or drawings which have been amended and are the basis for this report and/or sheets containing rectifications made before this Authority (see Rule 70.16 and Section 607 of the Administrative Instructions under the PCT).								
	These an	nexes consist of a total o	of 3 sheets.						
 - 		Basis of the opinion Priority Non-establishment of c Lack of unity of invention Reasoned statement in	on nder Rule 66.2(a)(ii) with regan ons supporting such statement ed	inventive step and industrial applicability rd to novelty, inventive step or industrial applicability;					
١	VIII Certain observations on the international application								
				·					
Date of submission of the demand				f completion of this report					
04.05.2004				3.2004					
Name and mailing address of the international preliminary examining authority: European Patent Office - Gitschiner Str. 103 D-10958 Berlin Tel. +49 30 25901 - 0 Fax: +49 30 25901 - 840				ized Officer , J one No. +49 30 25901-332					

INTERNATIONAL PRELIMINARY EXAMINATION REPORT

International application No.

PCT/EP 03/11218

I. Basis	of the	report
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1. With regard to the **elements** of the international application (Replacement sheets which have been furnished to the receiving Office in response to an invitation under Article 14 are referred to in this report as "originally filed" and are not annexed to this report since they do not contain amendments (Rules 70.16 and 70.17)):

	De	scription, Pages						
	1-4	19	as originally filed					
	Cla	aims, Numbers						
	6 (1	part), 7-12	as originally filed					
		6, 6 (part)	received on 04.05.2004 with letter of 26.04.2004					
2.	Wit lan	With regard to the language , all the elements marked above were available or furnished to this Authority in the language in which the international application was filed, unless otherwise indicated under this item.						
	The	These elements were available or furnished to this Authority in the following language: , which is:						
		the language of a translation furnished for the purposes of the international search (under Rule 23.1(b)).						
	☐ the language of publication of the international application (under Rule 48.3(b)).							
			anslation furnished for the purposes of international preliminant examination (under					
3.	With regard to any nucleotide and/or amino acid sequence disclosed in the international applicatio international preliminary examination was carried out on the basis of the sequence listing:							
			rnational application in written form.					
		filed together with th	e international application in computer readable form.					
	 ☐ furnished subsequently to this Authority in written form. ☐ furnished subsequently to this Authority in computer readable form. 							
	☐ The statement that the information recorded in computer readable form is identical to the written sequence listing has been furnished.							
4.	The	amendments have r	esulted in the cancellation of:					
		the description,	pages:					
		the claims,	Nos.:					
		the drawings,	sheets:					
5.		This report has been been considered to g	established as if (some of) the amendments had not been made, since they have so beyond the disclosure as filed (Rule 70.2(c)).					
		(Any replacement sh report.)	eet containing such amendments must be referred to under item 1 and annexed to this					
6.	Add	litional observations, i	f necessary:					

INTERNATIONAL PRELIMINARY EXAMINATION REPORT

International application No.

PCT/EP 03/11218

- V. Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement
- 1. Statement

Novelty (N)

Yes: Claims

1-12

No: Claims

Inventive step (IS)

Yes: Claims

1-12

No: Claims

Industrial applicability (IA)

Yes: Claims

1-12

No: Claims

2. Citations and explanations

see separate sheet

EXAMINATION REPORT - SEPARATE SHEET

Re Item V

Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

D1: WO 96/17840 cited in the application

D2: WO 95/30651 cited in the application

D3: DE 43 19 887 = WO9429267 cited in the application

D4: EP-A-0 398 072 cited in the application

D5: US-A-3 236 889

1. Novelty

Document D1 discloses structurally very close compounds according to formula (I) as claimed, which have an alkylether group instead of a propargylether group in their structure (see D1, compounds n° 100, 104-108, 110-113, etc. and example 14, compounds 14.1-14.3).

Documents D2-D5 refer to structurally different fungicides, which do not have the rest -X-NH-C(O)- according to formula (I) as claimed.

Claims 1-12 meet the criteria of Art. 33 (2) PCT.

2. Inventive step

- 2.1 According to the application (see especially page 1, I. 7-11) the problem underlying the invention is the provision of compounds having an improved fungicidal activity if compared to the compounds of the prior art D1-D4 cited in the application.
- 2.2 D1 represents the closest prior art, since it refers to structurally very close compounds also useful as fungicides.
- 2.3 It is stressed that the application does not contain any comparative data showing that the abovementioned problem has actually been solved by the technical features of the claimed compounds.
- 2.4 Consequently the problem underlying the present application should be seen in the provision of a further fungicides.



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- 2.5 In view of the examples and the biological examples of page 49 it is credible that this problem has actually been solved by the technical features of claim 1.
- 2.6 The proposed solution is in view of the teaching of D1-D5 considered as surprising. The skilled person looking for further fungicides and starting from the closest prior art D1 (see especially p. 33, example 14, compounds 14.1, 14.2 or 14.3) would not have been led to replace the (m)ethoxy group with a propargyloxy group since there is no indication in D1 nor D2-D5 that the propargyloxy group and the alkoxy group are bioisosteric substituents.

What is claimed is:

1. A compound of formula I

$$R_{1} = \begin{array}{c} R_{2} \\ R_{3} \end{array} \qquad \begin{array}{c} O - R_{4} \\ R_{5} \\ R_{6} \end{array} \qquad \begin{array}{c} O - R_{4} \\ R_{8} \end{array} \qquad (1)$$

including the optical isomers thereof and mixtures of such isomers, wherein R₁ is hydrogen, optionally substituted alkyl, optionally substituted cycloalkyl or optionally substituted aryl;

R₂, R₃, R₅, R₆, and R₇ are each independently of each other hydrogen or optionally substituted alkyl;

R₄ is optionally substituted alkyl;

X is O or N-R7; and

R₈ is a group

R₉ is optionally substituted aryl or optionally substituted heteroaryl;

R₁₀ and R₁₁ are each independently hydrogen, optionally substituted alkyl, optionally substituted alkenyl or optionally substituted alkynyl;

R₁₂ is optionally substituted alkyl, optionally substituted cycloalkyl, optionally substituted aryl or optionally substituted heteroaryl;

R₁₃ is hydrogen or optionally substituted alkyl, alkenyl or alkynyl; and R₁₄ is optionally substituted alkyl or optionally substituted amino.

- 2. A compound according to claim 1 wherein R_{10} is hydrogen or alkyl, X is oxygen, R_{8} is -C(R₉R₁₀)-OR₁₁ and R₁₁ is hydrogen or alkynyl.
- 3. A compound according to claim 1 wherein X is oxygen, R₈ is -C(R₁₂R₁₃)NH-SO₂-R₁₄, and R₁₂ is alkyl or branched alkyl. RAPER ROOM
- 4. A compound of formula I according to any of claims 1 to 3, wherein

 R_1 is hydrogen, alkyl, cycloalkyl, phenyl or naphthyl; phenyl and naphthyl being optionally substituted by substituents selected from the group comprising alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkyl, phenyl and phenylalkyl, where all these groups may in turn be substituted by one or several halogens; alkoxy; alkenyloxy; alkynyloxy; alkoxy-alkyl; haloal-koxy; alkylthio; haloalkylthio; alkylsulfonyl; formyl; alkanoyl; hydroxy; halogen; cyano; nitro; amino; alkylamino; dialkylamino; carboxy; alkoxycarbonyl; alkenyloxycarbonyl; or alkynyloxycarbonyl; and R_4 is alkyl; and R_8 is a group $-C(R_9R_{10})-OR_{11}$, R_9 is aryl or heteroaryl, each optionally substituted by substituents selected from to group comprising alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkyl-alkyl, phenyl and phenylalkyl, where all these groups may be substituted by one or several halogens; alkoxy, alkenyloxy, alkynyloxy; alkoxy-alkyl; haloal-koxy; alkylthio; haloalkylthio; alkylsulfonyl; formyl; alkanoyl; hydroxy; halogen; cyano; nitro; amino; alkylamino; dialkylamino; carboxy; alkoxycarbonyl; alkenyloxycarbonyl and alkynyloxycarbonyl; and R_{11} is hydrogen; alkyl or alkynyl; or R_8 is a group $-C(R_{12}R_{13})NH-SO_2-R_{14}$, and R_{14} is alkyl or alkylamino.

- 5. A compound of formula I according to any of claims 1 to 4, wherein R_1 is hydrogen, C_1 - C_8 -alkyl, C_3 - C_8 -cycloalkyl; and R_2 , R_3 , R_5 and R_6 are hydrogen; and R_4 is C_1 - C_6 -alkyl; and R_9 is phenyl, naphthyl, 1,3-biphenyl or 1,4-biphenyl, each optionally substituted by one to three substituents selected from the group comprising C_1 - C_8 -alkyl, C_2 - C_8 -alkenyl, C_2 - C_8 -alkynyl, C_1 - C_8 -haloalkyl, C_1 - C_8 -alkoxy, C_1 - C_8 -haloalkoxy, C_1 - C_8 -alkylthio, C_1 - C_8 -haloalkylthio, C_1 - C_8 -alkylsulfonyl, halogen, cyano, nitro and C_1 - C_8 -alkoxycarbonyl; and R_{10} is hydrogen or C_1 - C_4 -alkyl; and R_{11} is hydrogen, C_1 - C_8 -alkynyl; or C_2 - C_8 -alkynyl; and R_{12} is C_1 - C_8 -alkyl, C_3 - C_6 -cycloalkyl, C_3 - C_8 -alkenyl, C_3 - C_8 -alkynyl; phenyl or benzyl wherein the phenyl and benzyl is optionally substituted by one to three substituents selected from the group comprising C_1 - C_8 -alkyl, C_2 - C_8 -alkenyl, C_2 - C_8 -alkynyl, C_1 - C_8 -haloalkyl, C_1 - C_8 -alkoxy, C_1 - C_8 -haloalkoxy, C_1 - C_8 -alkoxycarbonyl; and R_{13} is hydrogen or C_1 - C_8 -alkylsulfonyl, halogen, cyano, nitro and C_1 - C_8 -alkoxycarbonyl; and R_{13} is hydrogen or C_1 - C_8 -alkyl; and R_{14} is C_1 - C_8 -alkyl; C_1 - C_8 -monoalkylamino or C_1 - C_8 -dialkylamino.
- 6. A compound of formula I according to any of claims 1 to 5, wherein R_1 is hydrogen or C_1 - C_6 -alkyl, and R_2 , R_3 , R_5 and R_6 are hydrogen; and R_4 is methyl or ethyl; and R_9 is phenyl or naphthyl each optionally substituted by one to three substituents selected from the group comprising C_1 - C_6 -alkyl, C_1 - C_6 -haloalkyl, C_1 - C_6 -alkoxy, C_1 - C_6 -haloalkylthio, halogen, cyano, nitro and C_1 - C_6 -alkoxycarbonyl; and R_{10} and R_{13} are